abcam

Product datasheet

Recombinant Human PFKM protein ab199591

描述

产品名称
重组人PFKM蛋白

纯**度** > 95 % SDS-PAGE.

Determined by SEC-HPLC and reducing SDS-PAGE

 内毒素水平
 < 0.100 Eu/μg</td>

 表达系统
 HEK 293 cells

Accession P08237

蛋白长度 Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列 THEEHHAAKTLGIGKAIAVLTSGGDAQGMNAAVRAVVRVGIF

TGARVFFV

HEGYQGLVDGGDHIKEATWESVSMMLQLGGTVIGSARCKDFR

EREGRLRA

 ${\tt AYNLVKRGITNLCVIGGDGSLTGADTFRSEWSDLLSDLQKAG}$

KITDEEAT

KSSYLNIVGLVGSIDNDFCGTDMTIGTDSALHRIMEIVDAIT

TTAQSHQR

TFVLEVMGRHCGYLALVTSLSCGADWVFIPECPPDDDWEEHL

CRRLSETR

TRGSRLNIIIVAEGAIDKNGKPITSEDIKNLVVKRLGYDTRV

TVLGHVOR

GGTPSAFDRILGSRMGVEAVMALLEGTPDTPACVVSLSGNQA

VRLPLMEC

VQVTKDVTKAMDEKKFDEALKLRGRSFMNNWEVYKLLAHVRP

PVSKSGSH

TVAVMNVGAPAAGMNAAVRSTVRIGLIQGNRVLVVHDGFEGL

AKGQIEEA

GWSYVGGWTGQGGSKLGTKRTLPKKSFEQISANITKFNIQGL

VIIGGFEA

YTGGLELMEGRKQFDELCIPFVVIPATVSNNVPGSDFSVGAD

IALNIICI

TCDRIKQSAAGTKRRVFIIETMGGYCGYLATMAGLAAGADAA

YIFEEPFT

IRDLQANVEHLVQKMKTTVKRGLVLRNEKCNENYTTDFIFNL

1

YSEEGKGI

FDSRKNVLGHMQQGGSPTPFDRNFATKMGAKAMNWMSGKIKE

SYRNGRIF

ANTPDSGCVLGMRKRALVFQPVAELKDQTDFEHRIPKEQWWL

KLRPILKI

LAKYEIDLDTSDHAHLEHITRKRSGEAAVVDHHHHHH

预测分子量 86 kDa including tags

氨基酸 2 to 780

标签 His tag C-Terminus

额外的序列信息 This product is for the mature full length protein. The initiator methionine is not included.

技术指标

Our **Abpromise guarantee** covers the use of **ab199591** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应**用** SDS-PAGE

HPLC

形式 Liquid

补充说明 This product was previously labelled as Fructose 6 Phosphate Kinase

制备和贮存

稳定性和存储 Shipped on Dry Ice. Store at -80°C. Avoid freeze / thaw cycle.

pH: 6.9

Constituents: 0.87% Sodium chloride, 99% Phosphate Buffer

Supplied as a 0.2 µm filtered solution

常规信息

通路 Carbohydrate degradation; glycolysis; D-glyceraldehyde 3-phosphate and glycerone phosphate

from D-glucose: step 3/4.

疾病相关 Defects in PFKM are the cause of glycogen storage disease type 7 (GSD7) [MIM:232800]; also

known as Tarui disease. GSD7 is an autosomal recessive disorder characterized by exercise intolerance with associated nausea and vomiting. Short bursts of intense activity are particularly difficult. Severe muscle cramps and myoglobinuria develop after vigorous exercise. Most patients obtain a "second wind" when the onset of exercise is followed by a brief rest period. In time

patients adjust their activity level and are well compensated.

序列相似性 Belongs to the phosphofructokinase family. Two domains subfamily.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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